## **Nucleus Network Pty Ltd**

Location:

-27.448675785146172, 153.02983773645755

Local Services Offered:

Early Phase, Clinical trials site, Phase 1 unit, Trial Patient Recruitment, Completion of study documentation as per ICH GCP and contract. Studies in Healthy Volunteers

Details: In addition to the information contained within csv generated by this website, this PDF provides additional site information

Title	Nucleus Network Pty Ltd		
Contact Person	Ric Navarro, Cameron Johnson, Jeffery Wong		
Language	English		
Services Offered	Early Phase, Clinical trials site, Phase 1 unit, Trial		
	Patient Recruitment, Completion of study		
	documentation as per ICH GCP and contract		
Contact Title	Vice President, Marketing & Business		
	Development		
Contact First Name	Ric		
Contact Last Name	Navarro		
Contact Email	r.navarro@nucleusnetwork.com.au		
Contact Phone Number	61 431 658 476		
Website Address	https://www.nucleusnetwork.com/au/		
Street Name and Number	300C Herston Rd		
Other areas of expertise	Healthy Volunteers		
Is your Facility affiliated with a government	No		
agency or part of a government funded health			
service?			
Are there any notable factors relating to your	Healthy Volunteers		
Patient Population (e			
Average time to start study (in calendar days)	Less than 30		
Does your Facility perform HREC	Yes		
(IRB/ERB/Ethics) Committee submissions?			
Does your Facility have a dedicated department	Yes		
or group to perform HREC			
(IRB/ERB/ETHICS) Committee submissions?			
Department contact name	Elaine Wong		
Department phone number	0402329162		
Department email address	e.wong@nucleusnetwork.com.au		
Initiating study activities approval	No		
Types of HREC (IRB/ERB/ETHICS)	Local		
Committee that are used			
Any Additional Process	No		

Does your Clinical Trial site or Service	Yes		
undertake any recruitment?			
Audit	Yes		
Audit Type	FDA (Food Drug Administration), Other		
Has your Clinical Trial Site or Service been	Yes		
accredited?			
If your Clinical Trial Site or Service has been	NATA		
accredited, please select all relevant types			
71			
Meeting frequency	Monthly		
Review required	Greater than 2 weeks		
Does the HREC require contract/budget	No		
approval prior to release of final approval			
documents?			
Does your Facility have other review boards	No		
that need to approve the study prior to HREC			
(IRB/ERB/Ethics) Committee submission? For			
example, scientific, radiation safety			
committees, or others			
If other review boards, please name	Radiation Safety Committee/Officer for clinical		
in other review boards, piedse name	trials needing radiology service and Independent		
	Expert Reviewer for FTIH studie		
Does your Facility have a written	Yes		
	l es		
SOP/Policy/Procedure for Informed Consent?			
Does your Facility have a written	No		
SOP/Policy/Procedure for Other vulnerable			
populations?			
Does your Facility have a written	No		
SOP/Policy/Procedure for Minor Assent for			
paediatric populations?			
Does your Facility have access to translators	No		
and translation support for study conduct (e.g.			
consent, study-specific instruction)?			
consont, study specific instruction):			
Does your Facility have a training program for	Yes		
the research staff?			
Does the course content include GCP?	Yes		
Do you have a process or program in place to	Yes		
retrain research staff when a protocol is			
amended?			
	1		

Does the study staff that prepares or transports dangerous goods have training that meets the IATA International Air Transport Association (US) or other countries hazardous training requirements for shipping dangerous goods?	Yes
Can your Facility support patient visits on weekends?	Yes
Can your Facility support in-patient admissions for research studies?	Yes
Is your Facility adequately staffed to support studies with both blinded and unblinded Investigational Product?	Yes
Does the Facility have storage space for Study-Related materials (e.g. Lab Kits, Patient Materials, etc.)?	Yes
Does your Facility have the ability to collect and store PK/PD specimens?	Yes
Does your Facility have the ability to collect PK/PD samples beyond normal business hours?	Yes
Does your Facility typically allow the collection of Pharmacogenomic (PGX) samples for research purposes?	Yes
Diagnostic equipment	CT Scan Computerized Tomography Scan, ECG/EKG Electrocardiogram, MRI Magnetic Resonance Imaging, X-Ray, Other
Additional equipment	TTE, EEG, Spirometry
Does your Facility have an SOP or process that ensures routine calibration and maintenance of general equipment? Examples of general equipment include: scale, pulse oximeter, stadiometer, sphygmomanometer, etc.?	
Does your Facility have the necessary equipment to treat medical emergencies (for example crash/code cart)?	Yes